

# 2-[(2-thio-3-ethyl-6-substitutedamino)-1,3,5-thiadiazino]imino-11-(piperazine-1-yl)dibenzo[b,f][1,4]oxazepines

D. T. Tayade<sup>1\*</sup>, P. R. Kale<sup>2</sup>

<sup>1</sup>Department of Chemistry, Government Vidarbha Institute of Science and Humanities, Amravati 444604.

<sup>2</sup>Department of Chemistry, S.R.R.Lahoti Science College, Morshi 444905.

\*Corresponding Author: D. T. Tayade

**Abstract:-** Recently a novel series of 2-[(2-thio-3-substituted-6-ethylamino)-1,3,5-thiadiazino] imino-11-(piperazine-1-yl) dibenzo[b,f][1,4] oxazepines [IXB(a-f)] was successfully synthesized by the isomerisation of 2-[(2-substitutedimino-6-ethylamino)-1,3,5-dithiozino] imino 11-(piperazine-1-yl) dibenzo [b,f][1,4] oxazepines [VIII(a-f)] by 5% aqueous sodium bicarbonate in ethanol medium. The structures of all synthesized compounds were justified on the basis of chemical characteristics, elemental analysis and spectral studies.

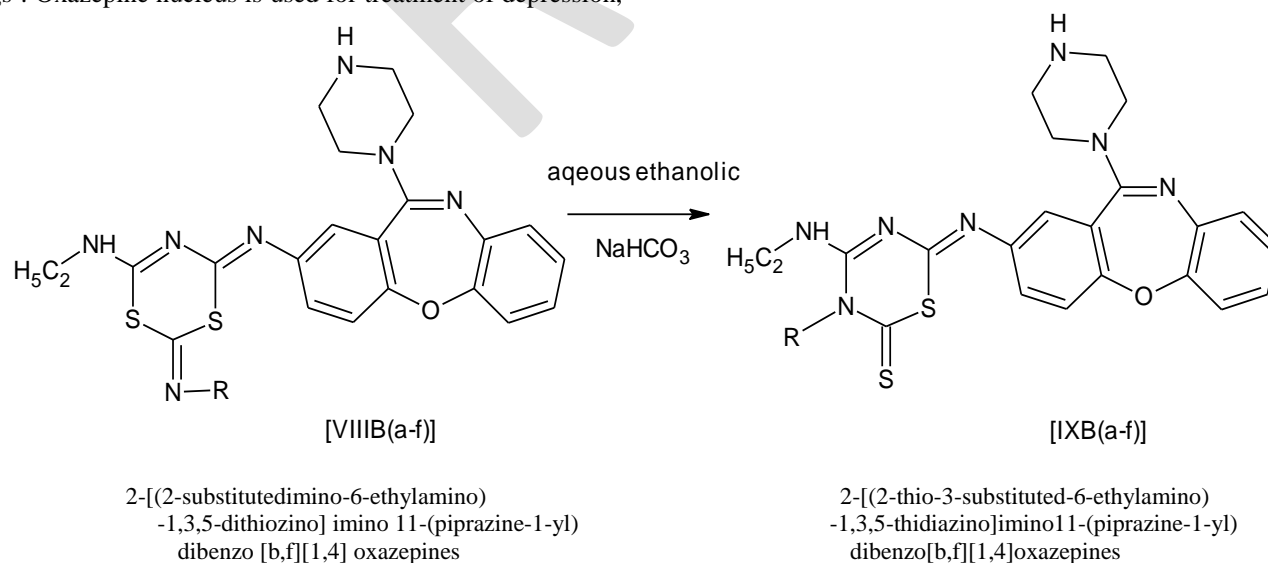
**Keywords:-** 2-[(2-substitutedimino-6-ethylamino)-1,3,5-dithiozino]imino11-(piperazine-1-yl)dibenzo[b,f][1,4] oxazepines and 5% aqueous sodium bicarbonate in ethanol.

## I. INTRODUCTION

Oxazepine and their derivatives have some important biological pharmacological activities<sup>1</sup> such as enzyme inhibitors<sup>2</sup>, analgesic<sup>3</sup>, anti-depressant<sup>4</sup> and psychoactive drugs<sup>5</sup>. Oxazepine nucleus is used for treatment of depression,

anxiety and agitation<sup>6-7</sup>. Recently new series of 1,2,4-thiadiazoles, 1,3,5-thiadiazines and 1,3,5-dithiazines were synthesized by exploring the synthetic applications of -thiocarbamido, -amino, -halo, -cyano, etc. and their antimicrobial, antifungal, antibacterial, analgesic physiochemical parameters<sup>8-11</sup> were studied. 2-Chloro-11-(piperazin-1-yl)dibenzo [b,f] [1,4] oxazepine (IB) and their derivatives showed agricultural, medicinal, biological, pharmaceutical, industrial significances and applications.

The main objective of the work is to synthesize a novel series of 2-[(2-ethylimino-6-substitutedamino)-1,3,5-dithiazino]imino-11-(piperazine-1-yl) dibenzo [b,f] [1,4] oxazepines [VIII(a-f)]. These were synthesized by the isomerisation of 2-[(2-substitutedimino-6-substitutedamino)-1,3,5-dithiozino]imino-11-(piperazine-1-yl) dibenzo [b,f][1,4] oxazepines [VIII(a-f)] by 5% aqueous sodium bicarbonate in ethanol, **Scheme-1**.

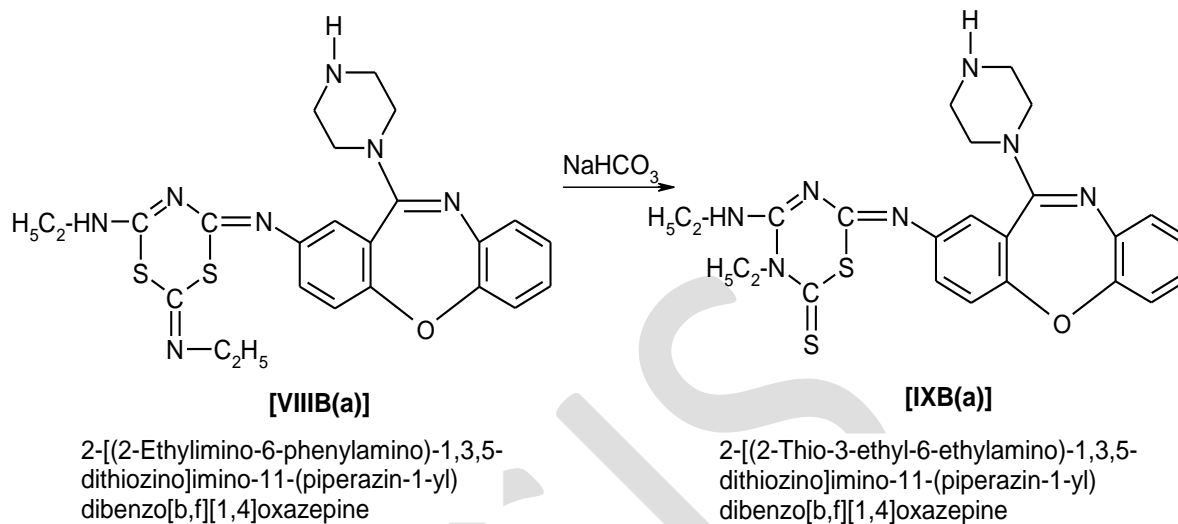


Where, R= -methyl, -ethyl, -t-butyl, -ethyl, p-chloroethyl, -p-tolyl.

**Scheme-1**

## II. SYNTHESIS OF 2-[(2-THIO-3-ETHYL-6-ETHYLAMINO)-1,3,5-THIAZINO]IMINO-11-(PIPERAZINE-1-YL)DIBENZO[B,F][1,4]OXAZEPINE

Synthesis of 2-[(2-thio-3-ethyl-6-ethylamino)-1,3,5-thiazino]imino-11-(piperazine-1-yl)dibenzo[b,f][1,4]oxazepine **[IXB(a)]** was carried out by isomerising 2-[(2-substituted imino-6-substitutedamino)-1,3,5-dithiozino]imino-11-(piperazine-1-yl)dibenzo [b,f][1,4]



**[IXB(a)]**

### III. PROPERTIES OF [IXB(a)]

It is brown colour crystalline solid having melting point 240°C. It gave positive test for nitrogen and sulphur. It was desulphurized by alkaline plumbite solution which clearly indicate the presence of C=S group. It was soluble in water, ethanol, DMSO-d<sub>6</sub> while insoluble in carbon tetrachloride, chloroform, benzene, petroleum ether. It formed picrate having melting point 209°C.

**Elemental analysis:** [C: 61.20% (found), 62.10% (calculated)], [H: 03.51% (found), 04.99 % (calculated)], [N: 18.11% (found), 18.11 % (calculated)], [S: 10.84% (found), 11.82 % (calculated)].

**IR Spectrum:** The IR spectrum was carried out in KBr-pellets. The important absorptions are correlated as (cm<sup>-1</sup>) 3180.62 N-H stretching, 2895.15 C-H stretching, 1726.89 N=C-N stretching, 1514.12 N-C=S stretching, 1288.45 C-N stretching, 1010.70 C=S stretching.

**NMR Spectrum:** The NMR spectrum was carried out in DMSO-d<sub>6</sub> and CDCl<sub>3</sub>. This spectrum distinctly displayed the signals due to Ar-H protons at δ 8.4227-6.9756 ppm, -NH proton at δ 3.6302-3.0104 ppm, -CH<sub>3</sub> protons at δ 1.3791-1.3437 ppm.

oxazepine **[VIIIb(a)]** in 5% aqueous sodium bicarbonate solution in ethanol. After distillation of excess solvent yellow crystals were separated out. Recrystallised from glacial acetic acid. Yield 94 %, m.p. 240°C.

The formation of **[IXB(b)]** is depicted below,

Similarly, 2-[(2-ethylimino-6-ethylamino)-1,3,5-dithiazino]imino-11-(piperazine-1-yl)dibenzo [b,f][1,4]oxazepine **[VIIIb(b)]**, 2-[(2-ethylimino-6-methylamino)-1,3,5-dithiazino]imino-11-(piperazine-1-yl)dibenzo [b,f][1,4]oxazepine **[VIIIb(c)]**, 2-[(2-ethylimino-6-tertbutylamino)-1,3,5-dithiazino]imino-11-(piperazine-1-yl)dibenzo [b,f][1,4]oxazepine **[VIIIb(d)]**, 2-[(2-ethylimino-6-p-chloroethylamino)-1,3,5-dithiazino]imino-11-(piperazine-1-yl)dibenzo[b,f][1,4]oxazepine **[VIIIb(e)]**, 2-[(2-ethylimino-6-p-tolylamino)-1,3,5-dithiazino]imino-11-(piperazine-1-yl)dibenzo [b,f][1,4]oxazepine **[VIIIb(f)]** were isomerized by 5% aqueous sodium bicarbonate solution by above mentioned method to isolate 2-[(2-thio-3-ethyl-6-ethylamino)-1,3,5-thiadiazino]imino-11-(piperazine-1-yl)dibenzo[b,f][1,4]oxazepine **[IXB(b)]**, 2-[(2-thio-3-ethyl-6-methylamino)-1,3,5-thiadiazino]imino-11-(piperazine-1-yl)dibenzo[b,f][1,4]oxazepine **[IXB(c)]**, 2-[(2-thio-3-ethyl-6-tertbutylamino)-1,3,5-thiadiazino]imino-11-(piperazine-1-yl)dibenzo[b,f][1,4]oxazepine **[IXB(d)]**, 2-[(2-thio-3-ethyl-6-p-chlorophenylamino)-1,3,5-thiadiazino]imino-11-(piperazine-1-yl)dibenzo [b,f][1,4]oxazepine **[IXB(e)]**, 2-[(2-thio-3-ethyl-6-p-tolylamino)-1,3,5-thiadiazino]imino-11-(piperazine-1-yl)dibenzo[b,f][1,4]oxazepine **[IXB(f)]**, by the above mentioned method and enlisted in **Table No. I**

Table No. I

Sr . N o.	Compd. No.	2-[(2-ethylimino-6- substitutedamino)1,3,5- dithiazino]imino-11-(piperazine- 1-yl) dibenzo [b,f] [1,4]oxazepine	Yield (%)	m.pt. (°C)
1	[IXB(b)]	2-[(2-Thio-3-ethyl-6- ethylamino)----- oxazepine	88	164
2	[IXB(c)]	2-[(2-Thio-3-ethyl-6- methylamino)----- oxazepine	92	148
3	[IXB(d)]	2-[(2-Thio-3-ethyl-6- butylamino)----- oxazepine	89	184
4	[IXB(e)]	2-[(2-Thio-3-ethyl-6- p-chlorophenylamino) ----- oxazepine	86	190
5	[IXB(f)]	2-[(2-Thio-3-ethyl-6- p-tolylamino)-----oxazepine	90	197

## REFERENCES

- [1]. Tripathi D.N., Malhotra RC. Bhattacharya A., J. Chromatography, 315 1984 417.
- [2]. Levinspial O., *Chem. Rea. Engg., John Willey and Sons* 2<sup>nd</sup> Ed, 1995.
- [3]. Aiello, F.; Brizzi, A.; Garofalo, A.; Grande, F.; Ragno, G.; Dayam, R.; Neamati, N. *Bioorg. Med. Chem.* 12, **2004**, 4459.
- [4]. Pecher J., Waefelaer A. *Bull. Soc. Chim. Belg.* **1978**, 87, 911.
- [5]. Halina K., Malgorzata S., Agata, W., 9(6), 828-850, 2012.
- [6]. Ayab H. *Journal of Al-Nahrain* 15 4 **2012** 47-59.
- [7]. J. Mikim, K. Y. Lee and J. N. kim. *Bull. Korean Chem.*; 23 (8), **2002** 1055-1056.
- [8]. Bansal R.K., *J.Heterocyclic Chemistry*, 8, **2012**, 12-24.
- [9]. Fernandes P.S and Sonar T.M., *J.Ind.Chem.Soc.*, 53(4),**1986**, 427.
- [10]. Saleem F., *Eur. Pat.*, CHAPPL 87/1 APR 13, 3600009 (1987), *Chem Abstr.*110,**1989**, 114893.
- [11]. Hedge J.C.,Satheesha Rai N. and Balkrishna K., *J.Chem.Sci.*,III 9(4),**2007**, 299-302.